

Administrator, Office of Diversion Control, hereby orders that the application submitted by the above firm for registration as a bulk manufacturer of the basic classes of controlled substances listed above is granted.

Dated: August 21, 1995.

Gene R. Haislip,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 95-21456 Filed 8-29-95; 8:45 am]

BILLING CODE 4410-09-M

Importer of Controlled Substances; Notice of Registration

By Notice dated July 5, 1995, and published in the **Federal Register** on July 13, 1995, (60 FR 36164), Applied Science Labs, Division of Alltech Associates, Inc., 2701 Carolean Industrial Drive, P.O. Box 440, State College, Pennsylvania 16801, made application to the Drug Enforcement Administration (DEA) to be registered as an importer of basic classes of controlled substances listed below:

Drug	Schedule
Heroin (9200)	I
Morphine (9300)	II

No comments or objections have been received. Therefore, pursuant to Section 1008(a) of the Controlled Substances Import and Export Act and in accordance with Title 21, Code of Federal Regulations, Section 1311.42, the above firm is granted registration as an importer of the basic classes of controlled substances listed above.

Dated: August 21, 1995.

Gene R. Haislip,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 95-21457 Filed 8-29-95; 8:45 am]

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Manufacturer of Controlled Substances; Notice of Application

Pursuant to Section 1301.43(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on July 12, 1995, Cambridge Isotope Lab, 50 Frontage Road, Andover, Massachusetts 01810, made application to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Cocaine (9041)	II

Drug	Schedule
Codeine (9050)	II
Methadone (9250)	II
Morphine (9300)	II

The firm plans to manufacture the controlled substances for isotope labeled standards for drug analysis.

Any other such applicant and any person who is presently registered with DEA to manufacture such substances may file comments or objections to the issuance of the above application.

Any such comments or objections may be addressed to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than October 30, 1995.

Dated: August 21, 1995.

Gene R. Haislip,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 95-21458 Filed 8-29-95; 8:45 am]

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Manufacturer of Controlled Substances; Notice of Application

Pursuant to Section 1301.43(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on July 17, 1995, Celgene Corporation, 7 Powder Horn Drive, Warren, New Jersey 07059, made application to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
2,5-Dimethoxyamphetamine (7396)	I
Amphetamine (1100)	II

The firm plans to manufacture small quantities of 2,5-Dimethoxyamphetamine using biocatalysis to develop, manufacture and sell high value added compounds to pharmaceutical and agrochemical industries and Amphetamine for distribution of the bulk active substance to its customers.

Any other such applicant and any person who is presently registered with DEA to manufacture such substances may file comments or objections to the issuance of the above application.

Any such comments or objections may be addressed to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement

Administration, United States Department of Justice, Washington, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than October 30, 1995.

Dated: August 21, 1995.

Gene R. Haislip,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 95-2145 Filed 8-29-95; 8:45 am]

BILLING CODE 4410-09-M

Federal Bureau of Investigation

DNA Advisory Board Meeting

Pursuant to the provisions of the Federal Advisory Committee Act, notice is hereby given that the DNA Advisory Board (DAB) will meet on September 20, 1995, from 9 a.m. until 5 p.m. at The Rockefeller University, Abby Aldrich Rockefeller Hall, Cohn Library, 1230 York Avenue, New York, NY 10021. All attendees will be admitted only after displaying personal identification which bears a photograph of the attendee.

The DAB's scope of authority is: To develop, and if appropriate, periodically revise, recommended standards for quality assurance to the Director of the FBI, including standards for testing the proficiency of forensic laboratories, and forensic analysts, in conducting analysis of DNA; To recommend standards to the Director of the FBI which specify criteria for quality assurance and proficiency tests to be applied to the various types of DNA analysis used by forensic laboratories, including statistical and population genetics issues affecting the evaluation of the frequency of occurrence of DNA profiles calculated from pertinent population database(s); To recommend standards for acceptance of DNA profiles in the FBI's Combined DNA Index System (CODIS) which take account of relevant privacy, law enforcement and technical issues; and, To make recommendations for a system for grading proficiency testing performance to determine whether a laboratory is performing acceptably.

The topics to be discussed at this meeting include: a review of minutes from the first two meetings; revised scope of authority for the DAB; a discussion of revisions to draft DAB by-laws; a review of the National Institute of Justice's (NIJ) DNA proficiency testing solicitation; an update on current forensic DNA technology; an overview of the FBI's Combined DNA Index System (CODIS); and a review of draft DNA testing standards based on current